DATA EVALUATION RECORD

SLUGKIL 5 (NEU 1182 M)

STUDY TYPE: ACUTE ORAL TOXICITY - RAT (870.1100) MRID 47942004

Prepared for Biopesticides and Pollution Prevention Division Office of Pesticide Programs U.S. Environmental Protection Agency One Potomac Yard 2777 South Crystal Drive Arlington, VA 22202

Prepared by Toxicology and Hazard Assessment Group **Environmental Sciences Division** Oak Ridge National Laboratory Oak Ridge, TN 37831 Task Order No. 10-006

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Disclaimer

Date:

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DATA EVALUATION RECORD

EPA Secondary Reviewer:

STUDY TYPE: Acute Oral Toxicity – Rats (OPPTS 870.1100)

MRID NO: 47942004

DP BARCODE NO: DP 373925

DECISION NO: 425380

SUBMISSION NO: 866102

TEST MATERIAL: NEU 1182 M (EPA Reg. No. 67702-GG)

PROJECT NO: 12326-08

SPONSOR: Neudorff GmbH KG, Postfach 1209, An der Mühle 3, D-

31860 Emmerthal, Germany

TESTING FACILITY: Stillmeadow, Inc., 12852 Park One Drive, Sugar Land,

TX 77478

TITLE OF REPORT: Acute Oral Toxicity

AUTHOR: Janice O. Kuhn

STUDY COMPLETED: November 19, 2008

GOOD LABORATORY GLP Compliant

PRACTICE:

CONCLUSION: The oral LD_{50} for female rats was greater than 5000

mg/kg.

CLASSIFICATION: ACCEPTABLE -- TOXICITY CATEGORY IV

I. STUDY DESIGN:

- 1. Test Material: NEU 1182 M, Chargen-Nr.: 070507
- 2. <u>Test Animals</u>: Three female Sprague-Dawley rats were received from Texas Animal Specialties, Humble, TX, and weighed 163-177 g on the day of dosing. The young adult animals, approximately 9 weeks old, were housed individually in suspended stainless steel cages with wire bottoms. The animals were fed PMI Feeds Inc. TM Formulab No. 5008, *ad libitum*. Municipal water was available *ad libitum*. The environmental conditions of the animal room were as follows: temperature, 22±3°C; relative humidity, 49-88% (outside protocol range but did not affect the outcome of the study); air changes, 10-12 per hour; and photoperiod, 12 hour light/dark cycle.
- 3. Methods: Rats were ear-punched: Nos. 1, 2, and 3, acclimated for 5 days, and fasted overnight prior to dosing. The test material was dosed at 5000 mg/kg body weight by gavage as a 40% w/v mixture in deionized water (Table 1). Body weight was recorded prior to dosing, and on days 7 and 14. The test animals were observed for mortality and clinical signs of toxicity at least three times on the day of dosing and daily thereafter for 14 days. All animals were necropsied.

II. RESULTS:

1. Mortality: All rats survived the study.

TABLE 1. Doses, mortality/animals treated			
Dose (mg/kg)	Males	Females	Combined
5000		0/3	

Data taken from p. 12, MRID 47942004.

- 2. Body Weight: All rats gained weight during the study.
- 3. Clinical Observations: All rats appeared normal throughout the study.
- 4. Gross Necropsy: No observable abnormalities were noted at necropsy.

III. DISCUSSION:

The oral LD₅₀ for female rats was greater than 5000 mg/kg. This places NEU 1182 M in TOXICITY CATEGORY IV. The packet classification is **ACCEPTABLE**.